

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
7 August 2003 (07.08.2003)

PCT

(10) International Publication Number
WO 03/063609 A1

(51) International Patent Classification⁷: A23K 1/14, C12P 7/06, A23J 1/12, 1/14, A23L 1/105, 1/211

(21) International Application Number: PCT/TB03/00289

(22) International Filing Date: 30 January 2003 (30.01.2003)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
BO2002A000047 30 January 2002 (30.01.2002) IT

(71) Applicant (for all designated States except US):
IBETECH S.R.L. [IT/IT]; Via San Basilio 44, I-45012
Ariano nel Polesine (IT).

(72) Inventors; and

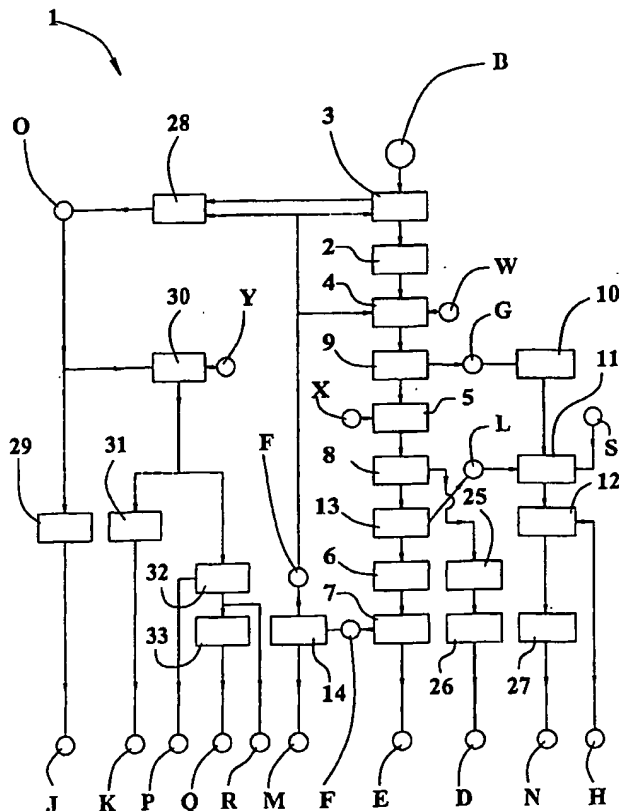
(75) Inventors/Applicants (for US only): ANGELINI, Andrea [IT/IT]; Via Alessandri 6, I-40126 Bologna (IT). MENEGATTO, Mario [IT/IT]; Strada Statale Romea 9, I-45019 Taglio di Po (IT). PEDRIELLI, Francesco [IT/IT]; Via M. M. Boiardo 2/D, I-44100 Ferrara (IT).

(74) Agent: AGAZZANI, Giampaolo; Agazzani & Associati S.r.l., Via Dell'Angelo Custode, 11/6, I-40141 Bologna (IT).

(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

[Continued on next page]

(54) Title: METHOD AND PLANT FOR CONVERTING GRAINS



(57) Abstract: A method for converting grains (B), containing at least starch, bran (D), germ (G) and proteins (L), includes a washing (3), a grinding (2), a first (4) and second (5) hydrolysis through enzymes, a fermentation (6) through yeasts and a distillation (7) of grains (B) to obtain alcohol (E) and a liquid. The method (1) provides: to subject grains (B) to solid fraction separation (8), by separating bran (D) before the fermentation (6); to subject to fermentation (6) grains (B) deprived of bran (D); to subject to distillation (7) grains (B), deprived of bran (D) and fermented, so obtaining alcohol (E) and an aqueous liquid (F) nearly without solid residues.

WO 03/063609 A1



(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

— before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

Published:

— with international search report

METHOD AND PLANT FOR CONVERTING GRAINS**TECHNICAL FIELD**

- 5 The present invention relates to vegetable products processing and it refers to a method and a plant for converting grains, particularly for using corn, soy, cereals and other similar agricultural products.

There are known methods and plants capable to obtain alcohols, such as ethanol, from
10 fermentation and distillation of corn grains and to generate a aqueous solution with high thermal contents and including, except the starch which is transformed by hydrolysis in glucose, practically all the different components present in the corn, particularly including a remarkable quantity of residual solid products a part of which, being soluble, passes in solution.

- 15 The main drawback of said known methods and plants consists in that the contained components in the aqueous solution, particularly proteins by-products, make said solution non reusable in advantageous way. Said solution, before being disposed, must be purified from the polluting components, particularly from solid residues, which must be separately disposed.

- 20 The thermal content of aqueous solution also represents a polluting element for disposal; in fact before proceeding to the disposal, it is necessary to provide also further energy expense for removing this thermal content from the aqueous solution.

Further drawback consists in that the presence of solid residues would make difficult the
25 thermal recycling of said thermal contents or of the solution water and therefore said known methods and plants require a constant and conspicuous supply of new water and of energy necessary to heating said water, causing an increase of the costs and of the quantity of polluting products to be disposed or dispersed in the environment.

- 30 There are further known methods and plants to obtain animal fodder based to grains in which the fodder in form of dry flour is added with enzymes and subjected to a hot extrusion process, up to over 200°C, that cooks the fodder, transforming it in a kind of compact and continuous paste, which then can be cut in pellets by a cutter, positioned at end of the extruder.

- 35 Said last known methods and plants have drawbacks consisting in that the excessive

temperature causes the inactivation of many enzymes, reducing the digestibility of the fodder; furthermore the introduction of "dry" state flour into the extruder requires high values of extrusion powers, causing high energetic cost and high usury of extruder components.

5

DISCLOSURE OF THE INVENTION

The main object of the present invention is to propose a method and a plant for converting grains for providing separately alcohol and an aqueous liquid almost without residual products,
10 in suspension or in solution, derived from the grain proteins.

Other object of the present invention is to propose a method and a plant for providing fodder, in pellets, flour or dry granular product, middle humidity or liquid fodders, having high digestibility and assimilation and safe for animal feeding, with very convenient energetic and
15 consumption costs, thanks to recoveries of energy and products provided by plant and method of the present invention.

Further object of the present invention is to propose a method and a plant fit for using the aqueous liquid and the related thermal contents and for providing bran, oil, amino acids and
20 yeasts, cereals and precooked and sanitized oily seeds, that is without foreign bodies, antinutritional factors, pathogenic bacteria, moulds and mycotoxins.

Other object is to propose a plant fit for effect a continuous cycle production.

25 Other object of the present invention is to propose a method and a plant capable to use the aqueous liquid and the related thermal contents for carrying out cereals, fermentable sugars, and therefore for transforming them in alcohol, or organic acids, antibiotic or other substances.

BRIEF DESCRIPTION OF THE DRAWINGS

30

The characteristics of the present invention are underlined in the following with particular reference to the attached drawings, in which:

- figure 1 shows a diagram of the method object of the present invention in which the phases are represented by rectangles and the initial, intermediate and final products are represented
35 by circles;

- figure 2 shows a schematic and partial view of the plant object of the present invention;
- figures from 3 to 6 show schematic views of details of figure 2 plant;
- figure 7 shows a schematic view of a variant of figure 2 plant.

5 BEST MODE OF CARRYING OUT THE INVENTION

With reference to figure 1, numeral 1 indicates the method for converting grain B, object of the present invention.

- 10 The grains B preferably consist of corn grains or soy, cereals or similar and mixtures thereof. It is provided that the grains can be added with materials derived from extraction of oil or others by-products.

The grains B have a fibrous portion, from now on indicated as bran D, and a germ G containing
15 an oily fraction H.

Furthermore grains B contain starch and proteins L in addition to several other substances such as vitamins, mineral salts and ashes.

- 20 The method 1 includes in sequence a washing 3, a grinding 2, a first hydrolysis 4 and a germ separation 9 fit for separating the germ G from the remaining portion of grains B.

The grains B without germ G are subjected to a second hydrolysis 5 and to separation of solid fraction 8, so obtaining bran D separated by the remaining part of grains B, which are thus
25 liquid and almost lacking of solid residues.

The grains B, without bran, are subjected to a protein separation 13, obtaining a protein fraction L separated from the remaining parts of grains B, lacking of protein fraction L.

- 30 The grains B, in liquid form, without germ, bran D, that is residual solid products, and without protein fraction L, are subjected to a fermentation 6 and to a distillation 7 obtaining separately alcohol E and a aqueous liquid F containing the less volatile fractions and almost lacking of residual products solid or in solution, except to fermentation yeasts.

- 35 Grinding 2 is quite rough in order to avoid excessive fragmentation of germ G.

Hydrolyses first 4 and second 5 are obtained through related enzymes mixed to grains B, brought at controlled temperatures, for instance around 95°C and 60°C respectively.

5. In correspondence of the hydrolyses first 4 and second 5, grains B are also mixed respectively with an alkalizing additive W, for solubilizing the proteins and an acidifier X for insolubilizing the proteins of grains B. Furthermore the alkalizing W and acidifying X additives provide the activation of respective enzymes.

10 The separations 9, 13 and 8 are carried out through forced decantation or similar separation processes.

Fermentation 6 is carried out by mixing yeasts to grains B, in the liquid form resulting from the preceding phases, and by maintaining the grains in a preset temperature interval for a predetermined time.

15 The distillation 7 is made by fractional evaporation of fermented grains.

The method provides the addition of water to the grains in correspondence of first hydrolysis 4 and of fermentation 6.

20 It is further provided that first and second hydrolyses can be unified in a single phase or that the total number of hydrolyses can be more than two.

25 Washing 3 and grinding 2 predispose grains B to the following manufacturing phases carrying out a mixture, in water, including bran D, starch, proteins L, oils H of germ G besides ashes, sugars and vitamins in proportions varying according to the nature of grains.

The first hydrolysis 4, thanks to the action of the related enzyme on the heated grains, converts the starch, which is splintered and adherent to the germ, in dextrins.

30 The second hydrolysis 5 converts the dextrins, deriving from the starch of the grains, in sugars, particularly in glucose, through the action of related enzyme and heating.

35 Fermentation 6 and distillation 7 carry out respectively the conversion of glucose in alcohol and the separation of the latter from the aqueous liquid F, containing yeasts and almost without

other solid residues, starches and glucose and having high thermal energy.

The method provides that the aqueous liquid F is eventually used, at least partially, for adding water to grains in method phases, for instance for the washing 3 and for contributing to the heating of grains B in correspondence of first hydrolysis 4.

The method 1 includes a yeast separation 14 from the aqueous liquid F to obtain yeasts M which developed during fermentation 6, separating them from the aqueous liquid F, which is thus purified, namely deprived of solids or of substances in solution. The yeasts M can be stored, dried, subject to hydrolysis and/or to autolysis, to breaking of the cellular membrane by mechanic or enzymatic means, wrapped, reused in the process or added to fodders.

It is also provided that grains B washed, prewashed by the washing 3 or by a further and separate washing phase, are subjected to cooking 28 in the purified aqueous liquid F obtaining sanitized grains B.

These sanitized grains B can be used as humid fodder or they can be subjected to desiccation 29, obtaining a dry fodder J, or they can be subjected to acidification and mixing 30 with solid and/or fluid alimentary additive Y, obtaining a semi finished product for fodder.

The semi finished product can be subjected to extrusion, cut and desiccation 31 obtaining fodder in pellets K or it can be subjected to homogenization 32 obtaining a homogenized fodder P, which can be directly used or subjected to spray desiccation 33 obtaining a soluble flour Q.

The method includes the thin grinding 10 of germ G obtained from the germ separation 9 and to subject the latter to proteolytic hydrolysis 11 together with the protein fraction L obtained from the proteins separation 13.

The proteolytic hydrolysis 11 resolves the germ membrane and it obtains the resolution of proteins released by the germ 13, together with the other proteins deriving from proteins separation 13, in amino acids thanks to the effect of a related enzyme activated by an additive S.

In addition the proteolytic hydrolysis 11 releases the oily portion of the germ.

The products of the proteolytic hydrolysis 11 are subjected to oil separation 12, by forced

decantation, obtaining, reciprocally separated, oils H and amino acids N in solution.

This last solution is subjected to concentration 27 obtaining a concentrated solution of amino acids N ready for fodder wrapping, storage or additive process.

5

The obtained oils H are delivered to suitable storage containers, ready for packaging and sale.

Bran D, separated from the remaining part of grains B by the solid fraction separation 8, is subjected, in sequence, to a desiccation 25 and to a bran grinding 26 obtaining bran D
10 containing gluten.

The plant 100 for converting grains, object of present invention, includes in cascade connection container and moving means 50 of grains B; washing means 51; grinding means 52; a first hydrolysis station 53; germ separating means 65, for instance of centrifugal type, provided with
15 an outlet for germ and an outlet for grains B without the germ.

The last outlet is connected in cascade to a second hydrolysis station 54 and to a separating station 55 provided with an outlet for a protein fraction, an outlet for a solid fraction, substantially consisting of bran, and an outlet for the remaining liquid fraction of grains B.
20

This last outlet is connected in cascade to a fermentation station 56 and to a distillation station 57, of evaporation and condensation type, having separate outlets for alcohol and for an aqueous liquid almost without solid fraction, starches, oils, and proteins.

25 Connecting means, including pipes, valves, pumping means and known and not shown intermediate reservoirs, connect the means and the stations of plant 100.

The container and moving means 50 of grains B include a collecting tank provided with a cochlea, known and not shown, for loading a silo container from which grains B are taken for
30 the transport, by a belt or the like, into washing means 51.

The grinding means 52 are capable to make a rough grinding, avoiding the pulverization of grains.

35 The grinding means 52 are connected to the first hydrolysis station 53 through hydration and

store means 63. These last means 63 are connected to water delivering means 61 for adding the grinded grains B with water and for compensating flow irregularity of grains B.

Each hydrolysis station first 53 and second 54 includes a thermoregulated enzymatic room 59 for grains B and into which enzyme delivering means 58 and additive delivering means 60 flow. The delivering means 58, 60 include respective tanks reservoirs and proportioners for regulating the quantities of enzymes and acidifying additive mixed with the grains B. Furthermore each hydrolysis station, first 53 and second 54, includes store means 62, consisting of tanks, positioned upstream and downstream the related enzymatic room 59 to allow an almost constant flow of grains B in the plant 100.

The first hydrolysis station 53 includes water delivery water 61 flowing into the respective enzymatic room 59 and heat exchanger means 64, for instance of countercurrent type, inserted into the inlet and outlet connections of the first hydrolysis station 53 for the heat regenerating.

It is also provided that the first 53 and second 54 hydrolysis stations can be integrated in a single hydrolysis station.

The fermentation station 56 is provided with fermentation yeast delivering means and with a connection, not shown, to the water delivering means 61.

The separation station 55 includes separating means of the solid fraction 68, of filter or centrifugal type, whose inlet is connected to the outlet of the second hydrolysis station 54 and whose outlets of liquid and solid fractions are respectively connected to an inlet of protein fraction separating means 67, of centrifugal type, and to bran treatment means 69.

The bran treatment means 69 include, in cascade, a warm air drier, a grindstone and a rotary feeder providing at the outlet bran D with gluten.

The protein fraction outlet of the protein fraction separating means 67 is connected to the second treatment means 70 and the non-protein fraction outlet is connected to the fermentation station 56.

The germ outlet of germ separating means 65 is connected to the second treatment means 70 through grinding means 71 capable to finely grind the germ fragments.

The second treatment means 70 include first store means 62 into which the protein fraction outlet of the protein fraction separating means 67 and the outlet of the grinding means 71 joint. The first store means 62 of second treatment means 70, feed proteolytic hydrolysis means 72 for the hydrolysis of the protein fraction and of the ground germ. The outlet of the proteolytic hydrolysis means 72 is connected, through second store means 62 of second treatment means 70, to amino acid separating means 73, whose amino acids outlet is connected to concentration means 74 for providing concentrated amino acids N. The remaining outlet of the amino acids separating means 73 provides an oily fraction H that is delivered to a respective containment reservoir for the further use.

10

It is also provided that the protein fraction separating means 67 and the solid fraction separating means 68 of the separating station 55 can be included in a single device of the type said tricanter.

15 The germ separating means 65, the separating station 55, the second treatment means 70 are connected to the water delivering means 61.

These last ones are connected to the waterworks or even to the aqueous liquid F outlet of distillation station 57.

20

This last outlet is connected to yeast separating means 75, of centrifugal type, an outlet of which provides yeasts M. The remaining outlet is connected to known cooking means 76 of grains B taken from washing means 51 to provides sanitized grains B by cooking at temperature lower than the temperature which causes a reduction of the alimentary values of sanitized and cooked grains.

25

It is also provided that suitable and specific washing means can feed the cooking means 76 with grains B.

30 The outlet of cooking means 76 is connected to third treatment means 78 cascade including, acidification means 79 and mixer means 80 fit for mixing grains sanitized and cooled after the cooking, with solid, fluid or containing enzymes alimentary additives Y. Furthermore the mixer means 80 can charge the sanitized grains B with amino acids N, and/or with other products of the plant 100, also transformed or modified.

35

The outlet of mixer means 80 is connected to pellet fodder production means 81, of known type, in order to provide pellet fodder K also in other commercial form.

It is also provided that the outlet of alimentary additive mixer means 80 can be connected to
5 flour fodder production means 82 including in cascade a homogenizer and a drier of so called spray type or having similar functions, to provide flour fodder P.

Furthermore it is provided that the outlet of cooking means 76 can be connected to desiccation means 77 to provide a dry fodder J.

10

Obviously the plant 100 can provide medium humidity or liquid fodders R.

The variant of plant 100 of figure 7 further include a leaching station 102 fed by oil-free grains Z, consisting of soy panels or residual seeds of an oil extraction process, and fed by the aqueous
15 liquid F, respectively through supply means 101 and through the distillation station 57.

The outlet of leaching station 102 is connected to sugar separating means 103, whose outlets separately provides a sugary fraction ZZ and a hyperproteic fraction LL of oil-free grains Z.

20 The outlet of this last fraction LL is connected to hyperproteic fraction concentration means 104 to provide the hyperproteic fraction LL in concentrated form.

The outlet of sugary fraction ZZ of sugars separating means 103 is cascade connected to alcoholic fermenter means 110 and to distiller means 111 to provide alcohol E.

25

Obviously, in alternative, the outlet of sugary fraction ZZ of sugar separating means 103 can be cascade connected to the fermentation station 56 and to the distillation station 57.

The outlet of the fraction sugary ZZ of the sugar separating means 103 can be also connected to
30 first 105 and second 106 fermenter means to provide respectively acids organic AO, antibiotics AN, which are concentrated by respective concentrators 112.

Furthermore, the outlet of sugary fraction ZZ of sugar separating means 103 can be connected to fermenter means, third 107 forth 108 and fifth 109, connected to respective concentrators 112
35 in order to obtain different substances for agricultural, zootechnical or industrial use, through

fermentations carried out by respective microorganism and/or mold strains.

It is provided that the outlet of sugar separating means 103 can be connected to respective proteolytic hydrolysis means 113 which feed a separator 114 fit for separately providing amino acids N and dietary fiber AF.

A possible operation of this variant provides to subject oil-free grains Z, mixes with aqueous liquid F, to leaching in acid environment and to separate the leaching product in a hyperproteic fraction LL, fit for concentration, and in its sugary fraction ZZ. The latter mainly consists of fermentable sugars, which are subjected to an alcoholic fermentation and to a distillation to obtain alcohol E.

Parallely or in alternative, the sugary fraction ZZ originates acids organic, antibiotics and other products through respective fermentations followed by related concentrations.

The main advantage of the present invention is to provide a method and a plant for converting grains for separately providing alcohol and an aqueous liquid practically without polluting solid residual products.

Other advantage of the present invention is to provide a method and a plant fit for providing fodders, in pellets, in flour form or in dry grains, medium humidity or liquid fodders, having high digestibility and assimilation and safe for animal feeding, said method and a plant being capable to use the aqueous liquid and the related thermal contents besides to provide bran, oil, amino acids and separate yeasts.

Other advantage is to provide a plant fit for effect a continuous cycle production.

Other advantage is to provide a method and a plant capable to use the aqueous liquid and the related thermal contents for carrying out from oil-free seeds fermentable sugars, and therefore for transforming them in alcohol or organic acids, antibiotics or other substances.

CLAIMS

- 1) Method for converting grains (B), containing at least starch, bran (D), germ (G) and proteins (L), including a washing (3), a grinding (2), at least a hydrolysis (4, 5) through enzymes, a fermentation (6) through yeasts and a distillation (7) of grains (B); said method (1) being characterized in that it provides:
- to subject grains (B), before fermentation (6), to a protein separation (13), obtaining a protein fraction (L) separated from grains (B);
 - to subject grains (B) without the protein fraction (L) to fermentation (6);
 - to subject to distillation (7) grains (B), without the protein fraction (L) and fermented, obtaining alcohol (E) and an aqueous liquid (F) practically without residual proteins by-products.
- 2) Method according to claim 1 characterized in that provides to include a first (4) and a second (5) hydrolysis and to subject grains (B) to solid fraction separation (8) downstream the second hydrolysis (5) obtaining bran (D) and bran-free grains (B) and to subject said grains to proteins separation (13) downstream of the solid fraction separation (8).
- 3) Method according to claim 1 characterized in that provides to subject the aqueous liquid (F) obtained by distillation (7) to yeast separation (14) achieving the purification of aqueous liquid (F) and obtaining yeasts (M).
- 4) Method according to claim 1 or claim 3 characterized in that provides to use at least partially the aqueous liquid (F) at least for washing (3).
- 5) Method according to claim 1 or claim 3 characterized in that provides to use at least partially the thermal energy of aqueous liquid (F) for heating grains (B) during at least one of the hydrolyses (4, 5).
- 6) Method according to claim 2 characterized in that provides to operate, in sequence, a desiccation (25) and a grinding (26) of the bran, obtaining bran (D) containing gluten, downstream the solid fraction separation (8).
- 7) Method according to claim 2 characterized in that provides:
- to effect a germ separation (9) from grains (B) after first hydrolysis (4) and before

fermentation (6);

- to subject the germ (G) to grinding (10);
- to subject the ground germ (G) to proteolytic hydrolysis (11) obtaining a solution containing an oily fraction (H).

5

8) Method according to claim 2 characterized in that provides:

- to charge grains (B) with at least an alkalizing additive (W) and acidifying additive (X) in correspondence of hydrolyses respectively first (4) and second (5);
- to subject the protein fraction (L) separated from grains (B) to proteolytic hydrolysis (11) obtaining amino acids (N) in solution;
- to subject the last solution to concentration (27) obtaining a concentrated solution of amino acids (N).

10

9) Method according to claims 7 and 8 characterized in that provides to subject the germ (G) to proteolytic hydrolysis (11) together with the protein fraction (L) downstream the grinding (10) of germ (G) and the proteins separation (13) and to subject the product of proteolytic hydrolysis (11) to oil separation (12) in order to separate the oily fraction (H) from the amino acids (N).

15

10) Method according to claim 1 characterized in that provides to take at least part of grains (B) from washing (3) and to subject said part to cooking (28) in the aqueous liquid (F) purified through yeast separation (14) obtaining sanitized grains (O).

20

11) Method according to claim 10 characterized in that provided to subject the sanitized grains (O) to desiccation (29) obtaining a dry fodder (J).

25

12) Method according to claim 10 characterized in that provides to subject the sanitized grains (O) to acidification and mixing (30) with alimentary additive solid and/or fluid (Y) obtaining a semifinished product for fodder.

30

13) Method according to claim 12 characterized in that provides to subject the semifinished product for fodder in sequence to extrusion, cut and desiccation (31) obtaining fodder in pellets (K).

14) Method according to claim 12 characterized in that provides to subject the semifinished

35

product for fodder to homogenization (32) obtaining a homogenized fodder (P).

15) Method according to claim 14 characterized in that provides to subject the homogenized fodder (P) to desiccation (33) obtaining a soluble flour (Q).

5

16) Method according to any of preceding claims characterized in that grains (B) consist of at least one among corn grains, soy, cereals, legumes or their mixtures.

17) Method according to any of preceding claims characterized in that first (4) and second (5) hydrolyses are integrated in a single hydrolysis phase.

10

18) Method according to any of preceding claims characterized in that provides:

- to mix oil-free grains (Z) with the aqueous liquid (F) and to subject the mixture to leaching in acid environment;
- 15 - to separate the leaching product in a hyperproteic fraction (LL), fit for concentration at least, and in a sugary fraction (ZZ);
- to subject sugary fraction (ZZ) to alcoholic fermentation and to distillation to obtain alcohol (E).

20 19) Method according to claim 18 characterized in that provides to subject the sugary fraction (ZZ) to fermentation and to a related concentration in order to obtain, separate and concentrated, at least one between organic acids (AO), antibiotics (AN).

20) Method according to claims 5, 7 and 8 characterized in that grains (B) to be subject to fermentation (6) substantially consist of a glucose solution carried out through the hydrolyses (4, 5) of the starch of grains (B) deprived, in sequence, of germ (G), bran (D) and proteins (L) respectively through germ separation (9), solid fraction separation (8) and proteins separation (13).

30 21) Plant for converting grains characterized in that it includes in cascade at least:

- container and moving means (50) of grains (B);
- washing means (51);
- grinding means (52);
- a first hydrolysis station (53);
- 35 - a second hydrolyses station (54);

- a separating station (55) having separate outlets at least for the solid fraction and the liquid fraction of grains (B);
- a fermentation station (56) connected to the liquid fraction outlet;
- a distillation station (57) having separate outlets for alcohol (E) and for a aqueous liquid (F).

5

22) Plant according to claim 21 characterized in that each of the first (53) and second (54) hydrolysis stations includes a thermoregulated enzymatic room (59) into which respective enzyme delivering means (58) flow.

10

23) Plant according to claim 22 characterized in that each enzymatic room (59) of first (53) and second (54) hydrolysis stations has additive delivering means (60).

24) Plant according to claim 22 characterized in that at least one of the hydrolysis stations, first (53) and second (54), includes water delivering means (61) flowing into the respective enzymatic room (59).

15

25) Plant according to claim 21 characterized in that each of the hydrolysis stations, first (53) and second (54), includes store means (62) positioned upstream and downstream the related enzymatic room (59).

20

26) Plant according to claim 21 characterized in that includes hydration and store means (63) interconnected between grinding means (52) and first hydrolysis station (53) and connected to water delivering means (61).

25

27) Plant according to claim 21 characterized in that at least one of the hydrolysis stations (53, 54) includes heat exchanger means (64) connected to the inlet and outlet connections of hydrolysis station (53, 54).

28) Plant according to claim 21 characterized in that the hydrolysis stations, first (53) and second (54), are integrated in a single hydrolysis station.

30

29) Plant according to claim 21 characterized in that the separating station (55) includes at least solid fraction separating means (68) whose inlet is connected to the outlet of the second hydrolysis station (54) and whose outlets of the liquid and solid fractions are connected

35

respectively to the inlet of fermentation station (56) and to bran treatment means (69).

30) Plant according to claim 29 characterized in that the liquid fraction outlet of solid fraction separating means (68) and the inlet of fermentation station (56) are connected through a non protein fraction inlet and outlet of protein fraction separating means (67) of the separating station (55), a protein fraction outlet of protein fraction separating means (67) is connected to second treatment means (70).

31) Plant according to claim 30 characterized in that includes germ separating means (65) having the inlet and the outlet for the germ-free fraction, interposed between the outlet and the inlet respectively of the hydrolysis stations first (53) and second (54), and having the germ outlet connected to second treatment means (70) through grinding means (71).

32) Plant according to claim 31 characterized in that the second treatment means (70) include proteolytic hydrolysis means (72) for the hydrolysis of the protein fraction and of the ground germ, whose outlet is connected to amino acids separating means (73), whose amino acids outlet is connected to concentration means (74), in order to provide concentrated amino acids (N), and whose remaining outlet provides an oily fraction (H).

33) Plant according to claim 30 characterized in that the protein fraction separating means (67) and the solid fraction separating means (68) are integrated in a tricanter device.

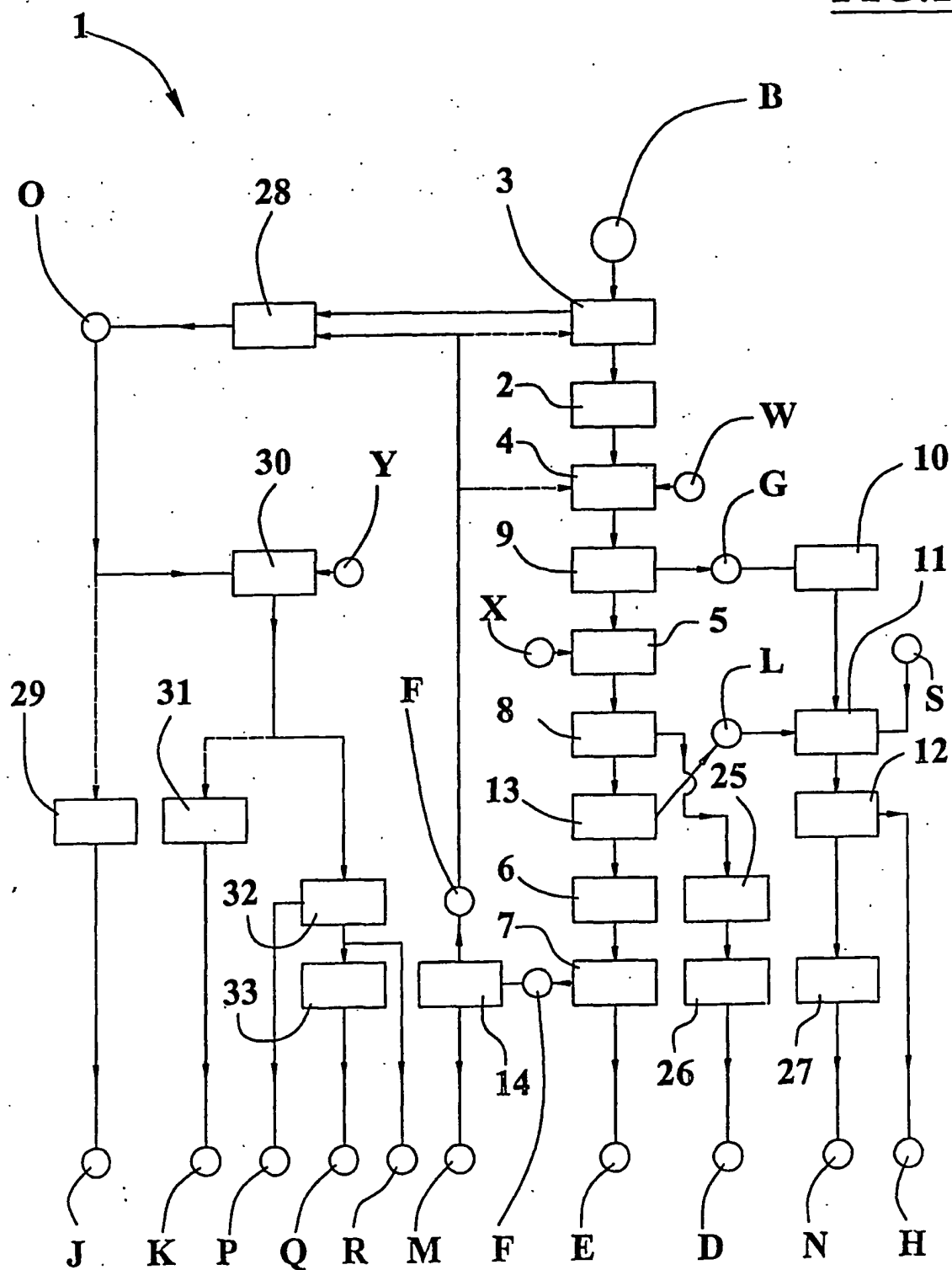
34) Plant according to claim 31 characterized in that germ separating means (65), separating station (55), second treatment means (70) are connected to water delivering means (61).

35) Plant according to claim 34 characterized in that the water delivering means (61) are connected to at least one between waterworks and aqueous liquid (F) outlet of the distillation station (57).

36) Plant according to claim 35 characterized in that the aqueous liquid (F) outlet of the distillation station (57) is connected to yeast separating means (75) an outlet of which provides yeasts and the remaining one feeds at least one between water delivering means (61) and cooking means (76) of grains (B), taken by washing means (51) in order to provide sanitized grains (B).

- 37) Plant according to claim 36 characterized in that the cooking means (76) outlet is connected to at least one between desiccation means (77), and third treatment means (78).
- 38) Plant according to claim 37 characterized in that the third treatment means (78) include, in cascade, acidification means (79) and alimentary additive mixer means (80) whose outlet is connected to at least one between pellets fodder production means (81) and flour fodder production means (82).
- 39) Plant according to any of claims from 21 to 38 characterized in that includes a leaching station (102), fed with free-oil grains (Z) and with aqueous liquid (F) respectively from supply means (101) and from the distillation station (57) and connected at the outlet to sugar separating means (103), whose outlets separately provide a sugary fraction (ZZ) and a fraction hyperproteic (LL) of free-oil grains (Z).
- 40) Plant according to claim 39 characterized in that includes fraction hyperproteic concentration means (104) to concentrate the fraction hyperproteic (LL).
- 41) Plant according to claim 39 characterized in that the sugary fraction (ZZ) outlet of sugar separating means (103) is cascade connected to alcoholic fermenter means (110) and to distiller means (111) in order to provide alcohol (E).
- 42) Plant according to claim 39 characterized in that the sugary fraction (ZZ) outlet of sugar separating means (103) is connected to at least one among first (105), second (106), third (107), forth (108), fifth (109) fermenter means to provide, at least, organic acids (AO) and antibiotics (AN).
- 43) Plant according to claim 42 characterized in that each outlet of the fermenter means (105 - 109) is connected to a respective concentrator (112).
- 44) Plant according to claim 39 characterized in that the outlet of sugar separating means (103) is connected to respective proteolytic hydrolysis means (113), which feed a separator (114) fit for separately providing amino acids (N) and dietary fiber (AF).

1/5

FIG.1

2/5

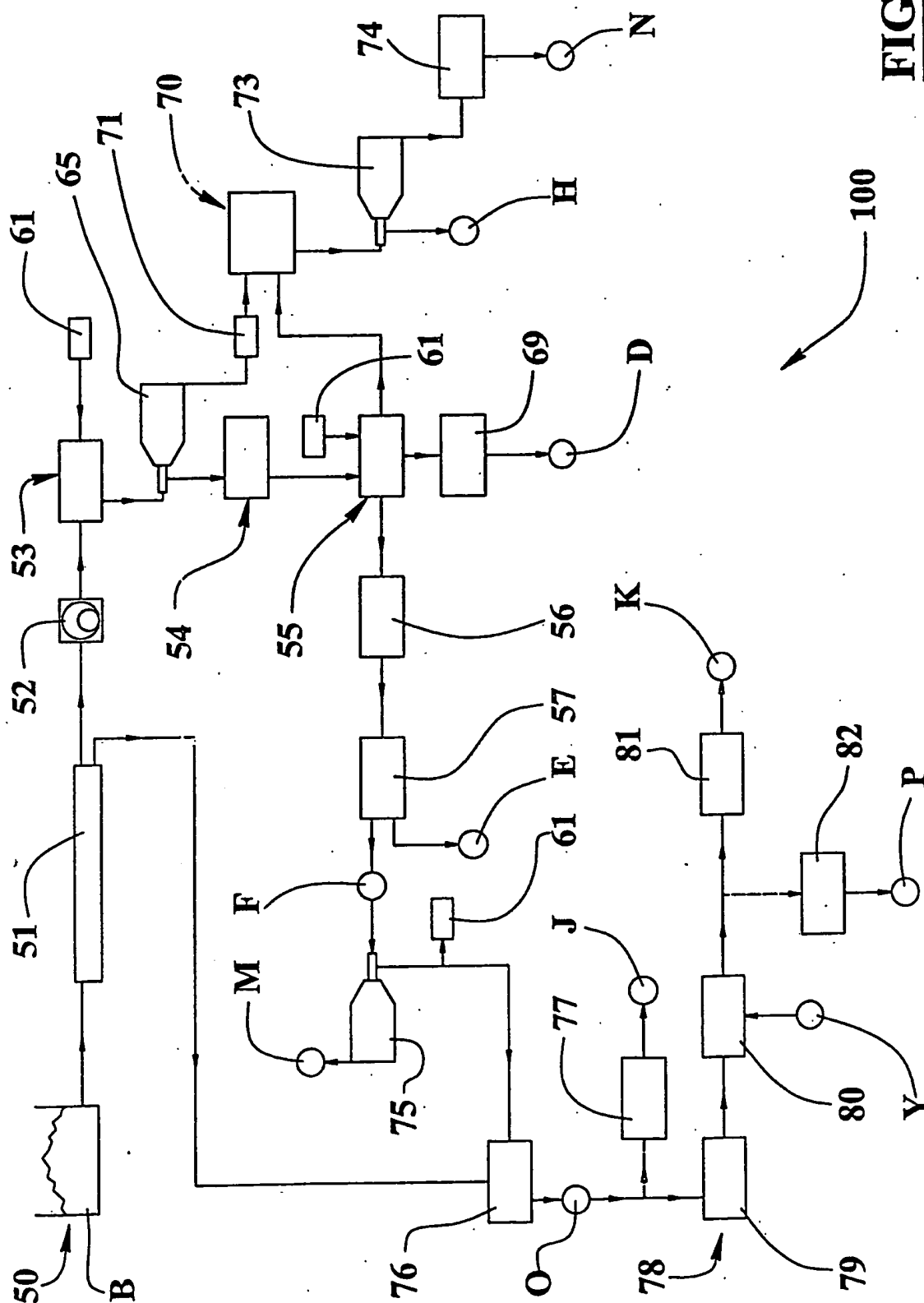


FIG.2

3/5

FIG.3

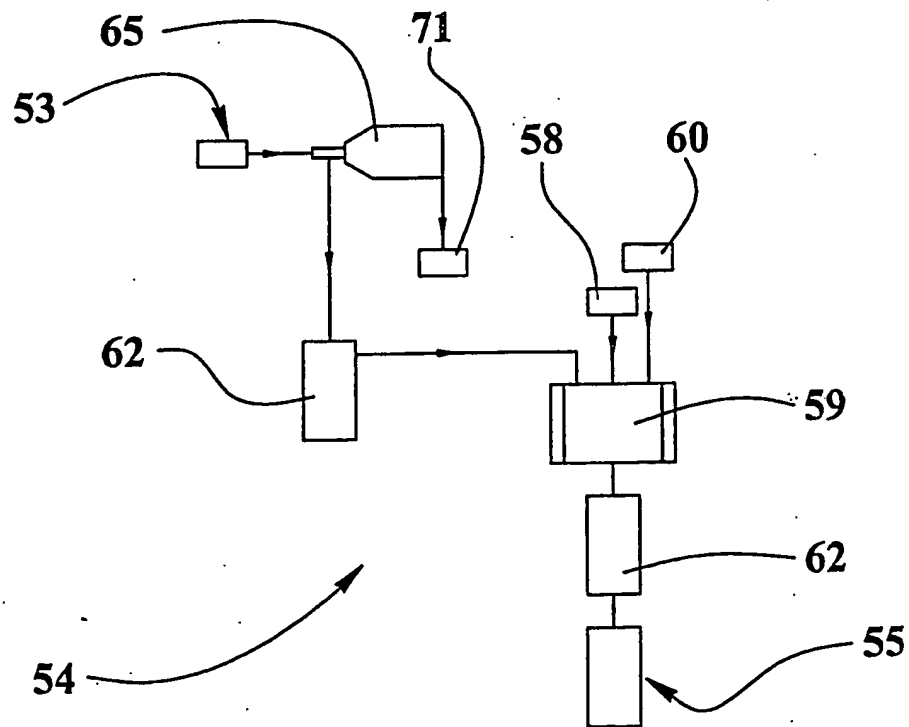
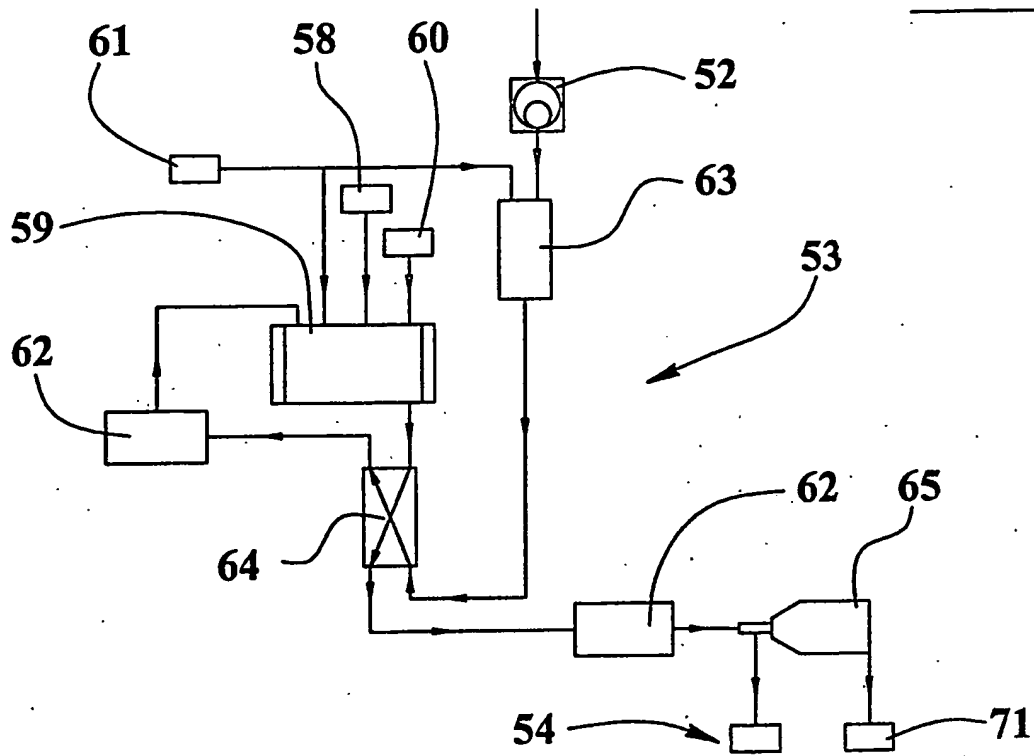


FIG.4

4/5

FIG.5

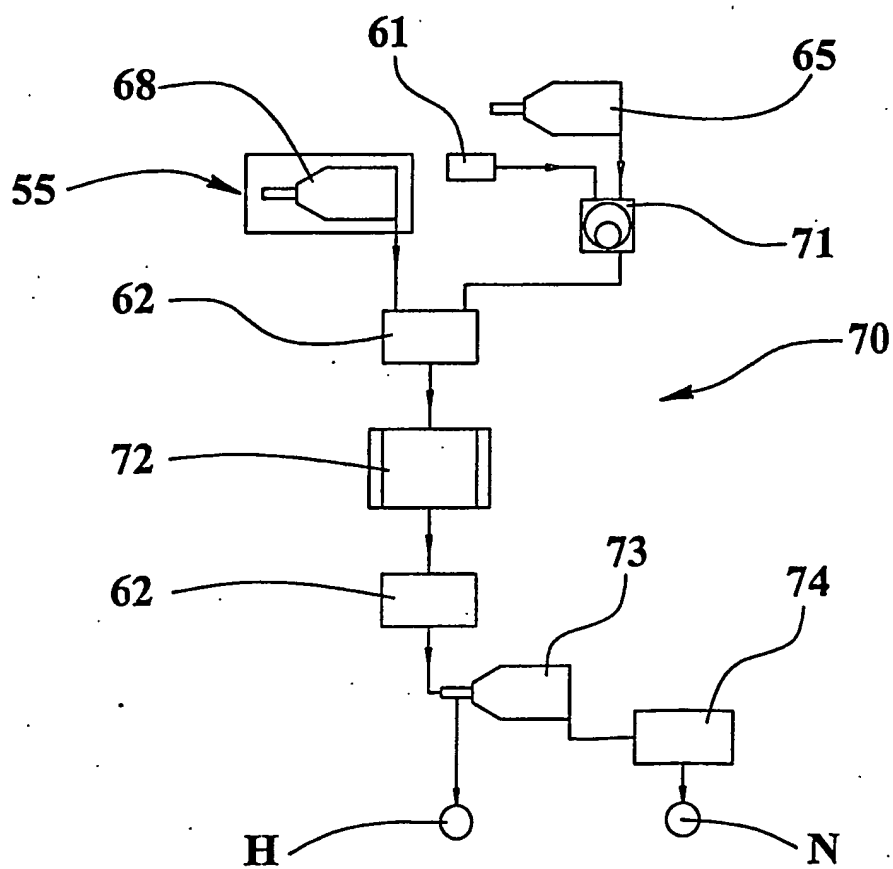
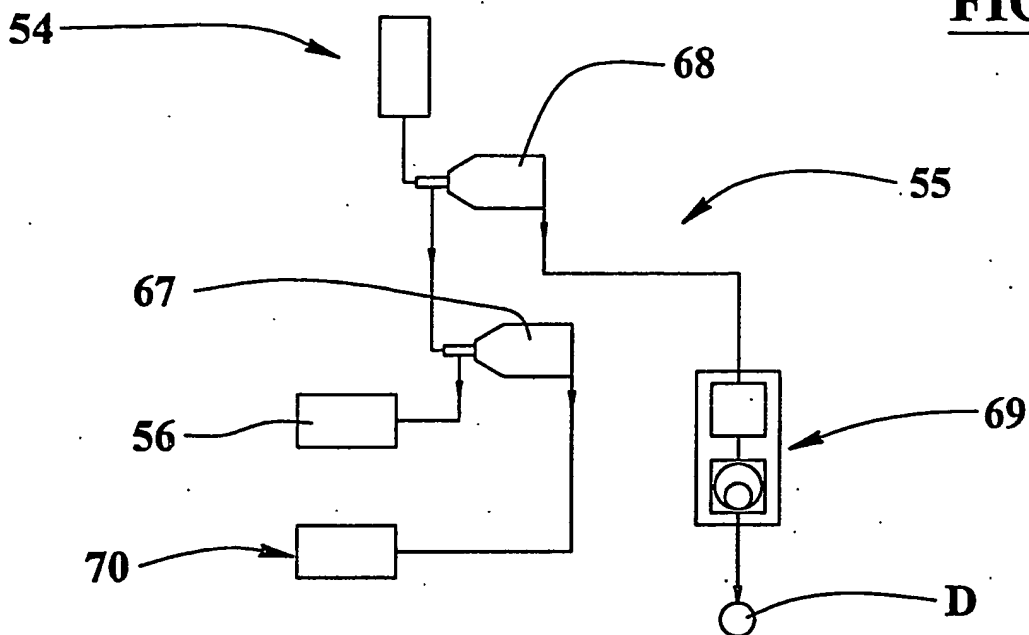
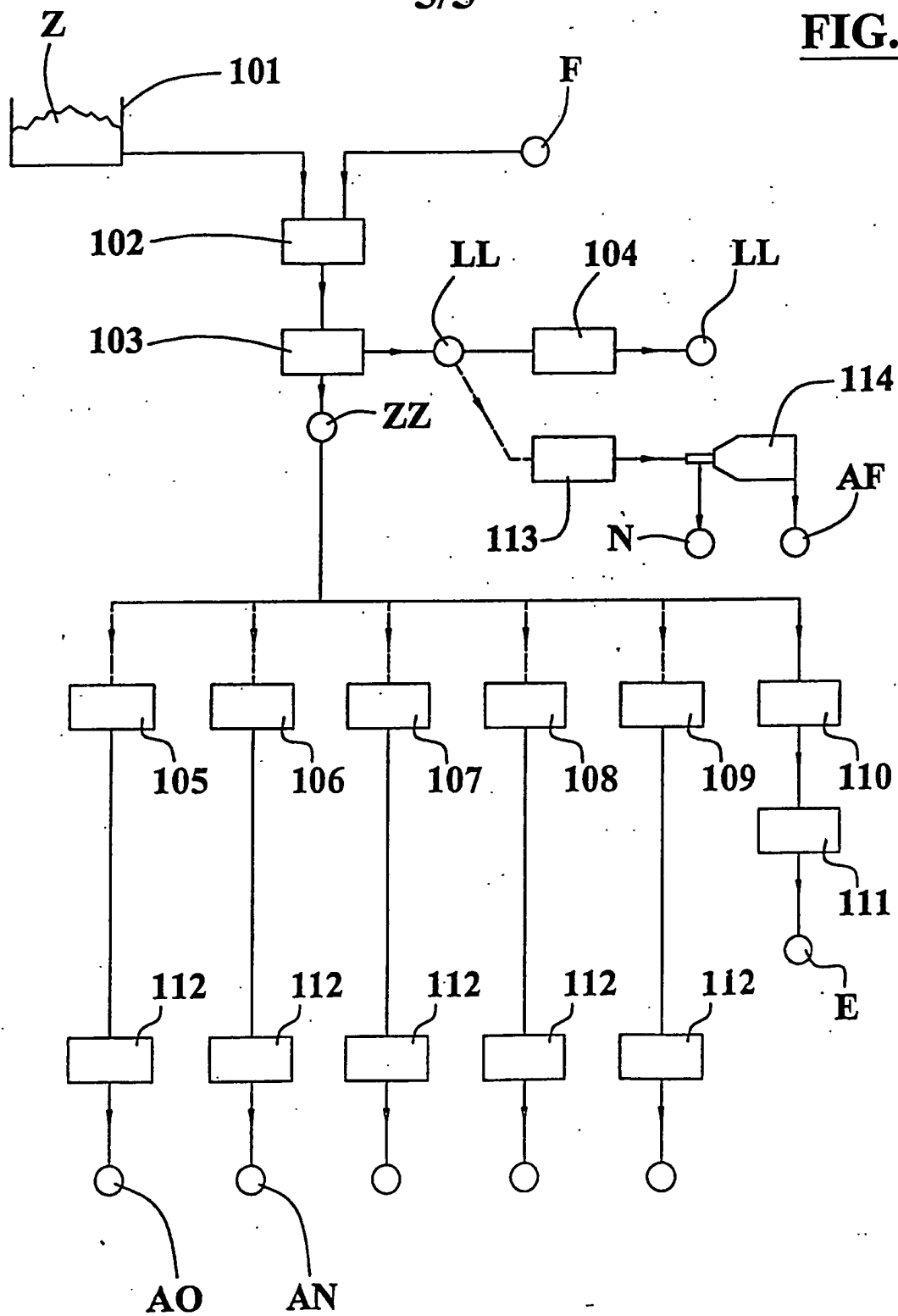


FIG.6

5/5

FIG.7

INTERNATIONAL SEARCH REPORT

Inte Application No

PCT/IB 03/00289

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A23K1/14 C12P7/06 A23J1/12 A23J1/14 A23L1/105
 A23L1/20 A23L1/211

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A23K C12P A23J A23L A23N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 4 448 881 A (MULLER WERNER C ET AL) 15 May 1984 (1984-05-15) the whole document ---	1-44
X	US 4 407 955 A (MULLER WERNER C ET AL) 4 October 1983 (1983-10-04) the whole document ---	1-44
X	AT 398 981 B (VOGELBUSCH GMBH) 27 February 1995 (1995-02-27) the whole document ---	1-44
A	US 4 810 647 A (MONCEAUX PHILIPPE ET AL) 7 March 1989 (1989-03-07) the whole document --- -/--	1-44

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the International filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the International filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- * & * document member of the same patent family

Date of the actual completion of the international search

15 May 2003

Date of mailing of the international search report

27/05/2003

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax: (+31-70) 340-3016

Authorized officer

De Jong, E

INTERNATIONAL SEARCH REPORT

Inte I Application No
PC 1 / 1 b 03/00289

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5 662 810 A (WILLGOHS RALPH H) 2 September 1997 (1997-09-02) column 1-2; claims 1-14 ---	1-44
A	WO 89 01522 A (TECHNIPETROL SPA) 23 February 1989 (1989-02-23) the whole document ---	1-44
A	GUTCHO M.H.: "Feeds for Livestock, Poultry and Pets" 1973 , NOYES DATA CORPORATION , LONDON XP002239398 page 105-108 ---	11-14, 37,38
A	US 4 001 452 A (WILLIAMS MERL A) 4 January 1977 (1977-01-04) the whole document ---	1-44
A	BE 1 003 192 A (DANIS) 7 January 1992 (1992-01-07) the whole document -----	1-44

INTERNATIONAL SEARCH REPORT

In International Application No

PCT/JP 03/00289

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 4448881	A	15-05-1984	US 4287304 A CA 1162157 A1	01-09-1981 14-02-1984
US 4407955	A	04-10-1983	NONE	
AT 398981	B	27-02-1995	AT 56991 A	15-07-1994
US 4810647	A	07-03-1989	FR 2586032 A1 BE 905227 A1 CA 1287003 A1 DE 3669530 D1 DK 372986 A EP 0213023 A1 ES 2001029 A6 IE 59029 B1	13-02-1987 04-02-1987 30-07-1991 19-04-1990 08-02-1987 04-03-1987 16-04-1988 15-12-1993
US 5662810	A	02-09-1997	US 5958233 A	28-09-1999
WO 8901522	A	23-02-1989	IT 1211714 B AT 133203 T AU 2260688 A DE 3854918 D1 EP 0330686 A1 WO 8901522 A1 US 5559031 A US 5545543 A	03-11-1989 15-02-1996 09-03-1989 29-02-1996 06-09-1989 23-02-1989 24-09-1996 13-08-1996
US 4001452	A	04-01-1977	BE 846075 A1 BR 7605887 A CA 1064759 A1 DE 2638292 A1 FR 2323340 A1 GB 1548703 A IT 1069570 B JP 52038375 A	31-12-1976 16-08-1977 23-10-1979 17-03-1977 08-04-1977 18-07-1979 25-03-1985 24-03-1977
BE 1003192	A	07-01-1992	BE 1003192 A6	07-01-1992